

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE
Examiner Lyle A. Alexander
ART UNIT 1743

In re application of
E. Alan Bates et al.
Application No. 08/935,629
Filed 09/23/97
For ASSAYING DEVICE

DECLARATION UNDER 37 CFR 1.132

Commissioner for Patents
Washington, DC 20231

I, Gary Hoffman, declare as follows:

1. I am the third joint inventor for this application.
2. While the claims of this application have been rejected based on the patent of Senior, the disclosure of the application starts from a technology actually quite different from that of Senior. Characteristic of this difference is Senior's provision of its bibulous member 16 protruding out of the housing. When the bibulous member is placed in a urine stream, it is quickly soaked. Senior has correctly chosen to display this version in its drawings, because versions such as suggested in paragraph a. in col. 5 of Senior would tend to shield the bibulous member from a urine stream.
3. This application references in its BACKGROUND a quite different technology, that of drug testing using an immunoassay method called antigen-antibody competitive binding. Characteristic of this technology is the measured application of only a few drops of urine. This dropwise application of the urine is disclosed in the specification in the first paragraph of the DETAILED DESCRIPTION OF THE INVENTION.
4. Attached hereto are Exhibits 1 to 5 demonstrating five different instances of the type

of technology forming the starting point for this application. These are as follows:

Exhibit 1 - Pages 1-4 of a document entitled "AccuSign DOA 4, THC/OPI/COC/AMP" bearing copyright notice dated 1996;

Exhibit 2 - Pages 1-4 of a document entitled "AccuSign BAR", also bearing copyright notice dated 1996;

Exhibit 3 - Front and back of a leaflet headed "Drug Test Resources International", likewise concerning AccuSign DOA 4 and bearing a 1996 copyright notice;

Exhibit 4 - One-sided, undated leaflet headed "Visaline II"; and

Exhibit 5 - Copies of the packages of several kits labeled HOME DRUG TEST and their instruction leaflets.

5. Exhibit 1 is noteworthy for the correspondence of the terminology in its section Principle on its page 1 with the terminology in the BACKGROUND section of this application.

6. All of the exhibits direct that 3 drops of urine be applied, this being a standard for this technology. All of the exhibits provide either a dropper or a pipette for transfer of the urine sample into the sample well. In each of the exhibits, the sample receiving area is clearly a well, and Exhibit 1, for instance, calls it a "well" in the section Test Protocol on its page 2.

7. The chart at the top of page 2 of Exhibit 1, for instance, explains that appearance of a line for a particular drug in this starting technology is a negative indication, i.e. the drug is not present in amounts above the cutoff level.

8. A characteristic of this technology is that the sample collection location must not be flooded with urine, as in Senior, because this leads to false positives by washing out the lines that would otherwise indicate negative readings.

FROM : HOFFMAN

FAX NO. : 412-821-2420

Aug. 21 2000 02:40PM P1

9. It is to be noted that none of these exhibits mentions photocopying the results. The statements regarding photocopying in the BACKGROUND section of the present application represent perceptions of the present inventor group, rather than the state of the art at the time this invention was made.

10. While the provision of a cap to cover Senior's wet, protruding member 16 is immediately understandable, it was the present group of inventors which perceived the advantages of such for the different technology from which the present invention arose.

11. A difference between the present invention and Senior concerns the problem with putting a cap on Senior's test. Because the bibulous material can be so flexible (especially when wet) there often is difficulty slipping the cap on without bending the material. It's like threading a needle by holding the thread still and moving the needle.

12. All statements made herein of my own knowledge are true and all statements made on information and belief are believed to be true; such statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and may jeopardize the validity of the application or any patent issued thereon.

Signature:

 Date: 8-21-00

AccuSign™ DOA 4

THC/OPI/COC/AMP

One-Step Panel Assay for Drugs of Abuse

For *In Vitro* Use Only

Simple One-Step Immunoassay for the Qualitative Detection of THC metabolites, Opiates, Cocaine metabolite, Amphetamines, and/or their Metabolites in Urine.

PBM

Catalog No.	DOA-240	35 Test Kit
	DOA-240-10	10 Test Kit

Intended Use

The AccuSign™ DOA 4 THC/OPI/CO/C/AMP Panel Assay is a simple, one-step immunochromatographic test for the rapid, qualitative detection of THC metabolites, opiates, cocaine metabolite, and amphetamines in urine. The test detects the major metabolites of these drugs at the following cutoff concentrations.

THC	11-nor-Δ ⁹ -THC-9-carboxylic acid	50 ng/mL
OPI	Morphine	300 ng/mL
COC	Benzoylecgone	300 ng/mL
AMP	Amphetamine	1000 ng/mL

The AccuSign™ DOA 4 THC/OPI/CO/C/AMP test provides only a preliminary analytical result. A more specific alternative chemical method must be used in order to obtain a confirmed analytical result. Gas chromatography, mass spectrometry (GC/MS) is the preferred confirmatory method. Other chemical confirmatory methods are available. Clinical consideration and professional judgment should be applied to any drug of abuse test result, particularly when preliminary positive results are used.¹

Summary and Explanation

Drug abuse has become one of the most destructive social problems in recent years, affecting nearly every corner of the world. To effectively combat this increasingly disturbing problem, there is a strong need for a simple, rapid, inexpensive, disposable, visual, and non-instrument requiring drug screening test kit. According to the National Institute on Drug Abuse (NIDA), THC (Marijuana), Opiates, Cocaine, and Amphetamines are among the most widely abused drugs. The one-step AccuSign™ DOA 4 Panel Assay is a test for screening these four major drugs of abuse in urine, simultaneously with one sample application. The test takes less than 10 minutes to perform.

THC (Δ⁹-tetrahydrocannabinol) is the primary active ingredient in cannabinoids (marijuana). When ingested or smoked, it produces euphoric effects. Users experience impairment of short term memory and THC use slows learning. Also, it may cause transient episodes of confusion, anxiety, or frank toxic delirium. Long term, relatively heavy use may be associated with behavioral disorders. The peak

effect of smoking THC occurs in 20–30 minutes and the duration is 90–120 minutes after one cigarette. Elevated levels of urinary metabolites are found within hours of exposure and remain detectable for 3–10 days after smoking. The main metabolite excreted in the urine is 11-nor-Δ⁹-tetrahydrocannabinol-9-carboxylic acid.

Opioid analgesics comprise a large group of substances which control pain by depressing the central nervous system. Morphine is the prototype compound of this group.² Morphine is excreted unmetabolized, and is also the major metabolic product of codeine and heroin. Morphine is detectable in the urine for several days after an opiate dose.

Cocaine, derived from the leaves of coca plant, is a potent central nervous system (CNS) stimulant and a local anesthetic. Cocaine induces euphoria, confidence and a sense of increased energy in the user; these psychological effects are accompanied by increased heart rate, dilation of the pupils, fever, tremors and sweating. Cocaine is used by smoking, intravenous, intranasal or oral administration, and excreted in the urine primarily as benzoylecgone in a short time. Benzoylecgone has a longer biological half-life (5–8 hours) than cocaine (0.5–1.5 hours) and can generally be detected for 24–60 hours after cocaine use or exposure.^{3,4}

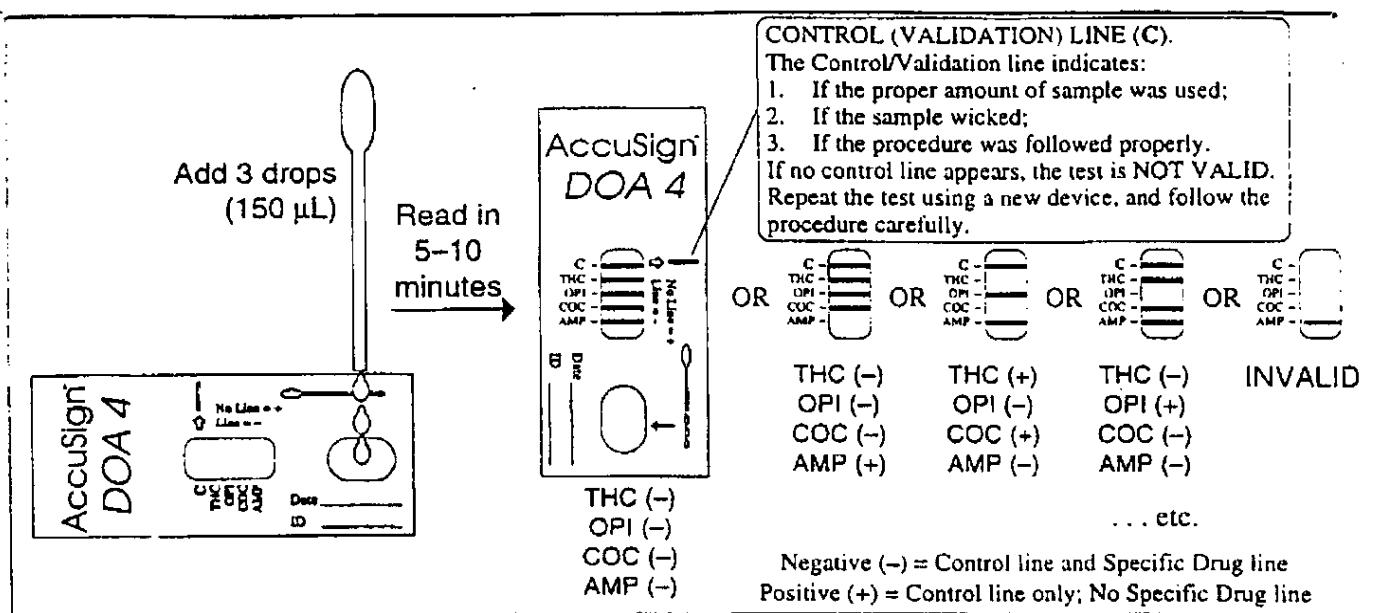
Amphetamine is a potent sympathomimetic agent with therapeutic applications. The drug can be taken orally, injected, or inhaled. Acute higher doses lead to enhanced stimulation of the central nervous system and induce euphoria, alertness, reduced appetite, and a sense of increased energy and power.⁵ Cardiovascular responses to amphetamine include increased blood pressure and cardiac arrhythmias. More acute responses include anxiety, paranoia, hallucinations, psychotic behavior, and eventually, depression and exhaustion. The effects of amphetamine generally last 2–4 hours, and the drug has a half-life of 9–24 hours in the body. Amphetamine is excreted in the urine in unchanged form and also as hydroxylated and deaminated derivatives.⁶

Principle

The AccuSign™ DOA 4 test employs one-step, solid-phase immunoassay technology to discretely detect the presence of any of the above four drugs, or their immunoreactive metabolites, in urine. The assay uses highly specific monoclonal/polyclonal antibodies raised against the target drugs. The test card contains a membrane strip, on which each of the four drugs conjugated to BSA is immobilized at specific locations. The assay is based on the principle of the highly specific immunochemical reactions between antigens and antibodies which are used for the analysis of specific substances in biological fluids. The drug detection relies on the competition for binding to the antibodies between drug conjugates and drugs which may be present in the urine sample.

In the test procedure, a sample of urine is placed in the sample well of the device, and the sample is allowed to migrate upward. If any of the four drugs is present in the urine sample, it forms a complex with the antibody-dye conjugate specific for that drug, and the complex migrates toward the opposite end of the card, passing the specific locations on the membrane where each of the four drug conjugates is immobilized. The drug in the sample competes with the drug conjugate, which is immobilized on the membrane, for the limited antibodies present in the form of antibody-dye conjugate. When a sufficient amount of drug is present, the drug will saturate the antibodies, and the antibody-dye conjugate cannot bind to the drug conjugate on the membrane. Therefore, a drug-positive urine sample will not generate a line at the specific drug position in the result window, indicating a positive result from positive drug competition. Conversely, if a particular drug is absent in the urine specimen, the antibody on the antibody-dye conjugate will bind the membrane-bound drug. In this case, a drug-negative urine sample will generate a line at the specific drug position in the result window, indicating a negative result from an absence of competition with free drug.

EXHIBIT



In addition, the test card has a procedural control built into the system, in the upper control line area. The control line is immobilized with polyclonal anti-mouse antibody; therefore, it will capture monoclonal antibody-dye conjugates that pass the region, showing a colored line in the control (validation) zone. The line works as a procedural control, confirming that proper sample volume was used and the reagent system worked. If insufficient sample volume is used, there may not be a control line, indicating the test is invalid.

Materials Provided

The AccuSign™ DOA 4 test kit contains all the reagents necessary to perform the assay.

- AccuSign™ DOA 4 device. The test device contains a membrane coated with drug conjugates in a protein matrix and a pad containing mouse monoclonal anti-THC antibody-dye conjugate, mouse monoclonal anti-opiate antibody-dye conjugate, mouse monoclonal anti-benzoylecgonine antibody-dye conjugate, and polyclonal sheep anti-amphetamine antibody-dye conjugate in a protein matrix.
- Disposable sample dispenser.
- Instructions for use.

Precautions

- For *in vitro* diagnostic use only.
- Avoid cross contamination of urine samples by using a new urine specimen container and dropper for each urine sample.
- This test kit does not contain any HIV or hepatitis infective components. However, urine specimens are potentially infectious. Proper handling and disposal methods should be followed, according to good laboratory practices.
- The AccuSign™ device should remain in its original sealed pouch until ready for use.
- Do not use the test kit after the expiration date.

Storage and Stability

The AccuSign™ DOA 4 test kit should be stored at 2-30°C (35-86°F) in the original sealed pouch. The expiration dating was established under these storage conditions.

Specimen Collection and Preparation

Approximately 150 µL of urine sample is required for each test. Fresh urine specimens do not require any special handling or pretreatment. Specimens should be collected in a clean glass or plastic container. If

testing will not be performed immediately, specimens should be refrigerated (2-8°C) or frozen. Specimens should be brought to room temperature before testing.

Specimens containing a large amount of particulate matter may give inconsistent test results. These specimens should be clarified by centrifuging or allowing to settle before testing.

Test Procedure

The test procedure consists of adding the urine sample to the Sample well of the device and watching for the appearance of colored lines in the result window.

Test Protocol

- For each test, open one AccuSign™ DOA 4 pouch and label the AccuSign™ device with the patient ID.
- Holding the dropper vertically, dispense 3 full drops (150 µL) of the urine sample into the Sample well.
- Read the result after 5-10 minutes.

Interpretation of Results

Negative: The appearance of a reddish-purple Control line (C) and a line for a specific drug indicates a negative test result; i.e., no drug above the cutoff level has been detected. The color intensities of the Control line and specific drug line may not be equal. *A negative test result does not indicate the absence of drug in the sample; it indicates only that the sample does not contain drug above the cutoff level in qualitative terms.*

Positive: The appearance of only a reddish-purple Control line and no distinct line next to a specific drug name indicates the test result is positive for that drug (i.e., the specimen contains the drug at a concentration above the cutoff level). *A positive test result does not provide any indication of the level of intoxication or urinary concentration of the drug in the sample; it indicates only that the sample contains drug above the cutoff level in qualitative terms.*

Invalid: A distinct Control line (C) should always appear. The test is invalid if no Control line forms at the C position. Such tests should be repeated with a new AccuSign™ DOA 4 test device.

Examples of possible results are shown in the diagram above.

- THC (-), Opiates (-), Cocaine (-), Amphetamines (-):** Five reddish-purple lines—one Control line at the C position and one each at the THC, OPI, COC, and AMP positions.
- THC (-), Opiates (-), Cocaine (-), Amphetamines (+):** Four reddish-purple lines—one Control line at the C position and one line each at the THC, OPI, and COC positions; no line at the AMP position.
- THC (+), Opiates (-), Cocaine (+), Amphetamines (-):** Three reddish-purple lines— one Control line at the C position, one line each at the OPI and AMP positions; no lines at the THC and COC positions.
- THC (-), Opiates (+), Cocaine (-), Amphetamines (-):** Four reddish-purple lines—one Control line at the C position and one line each at the THC, COC, and AMP positions; no line at the OPI position.
- There are other possible results, depending on the combinations of drugs present in the urine sample.

Note: A very faint line for a specific drug in the result window, visible in 10 minutes, indicates that the amount of drug in the sample is near or below the cutoff level of the test. These urine specimens must be retested, or confirmed with a more specific alternative method such as gas chromatography/mass spectrometry, before positive determinations are made.

Limitations

- The test is designed for use with unadulterated urine only.
- There is a possibility that factors such as technical or procedural errors, as well as other substances in the urine sample than those listed in Table 4 below, may interfere with the test and cause erroneous results.
- Adulterants, such as bleach and/or alum, in urine specimens may produce erroneous results regardless of the method of analysis. If adulteration is suspected, the test should be repeated with a new sample.
- This test detects only the presence of THC metabolites, opiates, cocaine metabolite, amphetamines, and/or their metabolites in urine. A positive test result does not provide any indication of the level of intoxication or urinary concentration.
- The test result read after 10 minutes may not be consistent with the original reading obtained within the 10 minute reading period. The test result must be read within 10 minutes of sample application.
- Certain medications containing opiates or opiate derivatives, amphetamines, or methamphetamines may produce a positive result in any chemical or immunological assay. Additionally, foods and tea containing poppy products and/or coca leaves may produce a positive result. Prolonged passive smoking of THC may also produce a positive result.

User Quality Control

Quality Control: Control standards are not supplied with this kit; however, it is recommended that a control be tested as good laboratory testing practice. NIDA recommends that positive quality control specimens be at or near the cutoff concentration. For information on how to obtain controls, contact PBM's Technical Services. Before using a new kit with patient specimens, positive and negative controls should be tested to confirm the test procedure, and to verify the tests produce the expected Q.C. results. Q.C. specimens should also be run anytime there is any question concerning the validity of results obtained.

Process Control: The Control line can be considered an internal process control. A distinct reddish-purple Control line should always appear if the test procedure is performed properly, an adequate sample volume is used, the sample and reagent are wicking on the membrane, and the test reagents are working. If the Control line does not appear

in the control or validation line area, the test is invalid and a new test should be performed. If the problem persists, contact PBM for technical assistance.

Expected Values

AccuSign™ DOA 4 is a qualitative assay. The amount of drugs and metabolites present in urine cannot be estimated by the assay. The assay results distinguish positive from negative samples. Positive results indicate the samples contain the specific drug above the cutoff concentration.

Performance Characteristics

The AccuSign™ DOA 4 Panel Assay detects THC, opiates, cocaine, amphetamines, and their metabolites at cutoff levels based on the recommendations of the National Institute on Drug Abuse (NIDA) for screening of these drugs in urine.^{13a}

THC	11-nor-Δ ⁹ -THC-9-carboxylic acid	50 ng/mL
OPI	Morphine	300 ng/mL
COC	Benzoylcegonine	300 ng/mL
AMP	D-Amphetamine	1000 ng/mL

The accuracy of AccuSign™ DOA 4 was evaluated in comparison to a commercially available immunoassay (Syva® EMIT® II) for each of these four drugs. About 1000 random clinical samples for each drug, including at least 250 positive samples above the cutoff level for each of the four drugs, was tested by both procedures, using the cutoff values listed. Complete agreement was observed in > 99% of the samples as shown below. (Table 1.)

Table 1. Accuracy: Comparison of AccuSign™ DOA 4 with Syva® EMIT® II Assay

Syva® EMIT® II (THC)			
	Positive	Negative	TOTAL
AccuSign™ Positive	305	5	310
DOA 4 (THC) Negative	11	680	691
TOTAL	316	685	1001

Syva® EMIT® II (OPI)			
	Positive	Negative	TOTAL
AccuSign™ Positive	249	0	249
DOA 4 (OPI) Negative	1	716	717
TOTAL	250	716	966

Syva® EMIT® II (COC)			
	Positive	Negative	TOTAL
AccuSign™ Positive	362	1	363
DOA 4 (COC) Negative	14	644	658
TOTAL	376	645	1021

Syva® EMIT® II (AMP/MET)			
	Positive	Negative	TOTAL
AccuSign™ Positive	185	0	185
DOA 4 (AMP) Negative	4	291	295
TOTAL	189	291	480

	Relative Sensitivity	Relative Specificity
THC	96.5% (305/316)	99.2% (680/685)
Opiates	99.6% (249/250)	> 99% (716/716)
Cocaine	96.3% (362/376)	99.8% (644/645)
Amphetamine	97.8% (185/189)	> 99% (291/291)

In a separate study, AccuSign™ DOA 4 was evaluated against specimens confirmed as positive by GC/MS, for each of the four drugs. The results below demonstrate the excellent correlation of AccuSign™ DOA 4 with GC/MS (99% agreement). (Table 2.)

Table 2. Accuracy: Comparison of AccuSign™ DOA 4 with GC/MS Assay

		AccuSign™	GC/MS
THC	Positive	87	88
	Negative	1	0
OPI	Positive	73	74
	Negative	1	0
COC	Positive	77	78
	Negative	1	0
AMP	Positive	55	56
	Negative	1	0

Precision and Accuracy

The precision of the AccuSign™ DOA 4 Panel Assay was determined by carrying out the test with serially diluted standard drug solutions. About 98% of the samples containing cocaine, opiates, or amphetamine and about 90% of the samples containing THC concentrations 25% over the cutoff level consistently showed positive results.

The study also included over 40 samples \pm 25% cutoff level as a challenge of cutoff precision. These results were found to be consistently in agreement with expected test results.

Distribution of Random Error:

Twenty (20) blind samples prepared by spiking various concentrations of cocaine, THC, morphine, or amphetamine were separately tested by two operators. The test results from the two operators showed complete agreement.

Reproducibility

The reproducibility of the test results of the AccuSign™ DOA 4 Panel Assay was examined at three different sites using a total of 15 blind controls, consisting of 5 negative samples, 5 moderately positive samples, and 5 strongly positive samples (i.e., a concentration 3 times the cutoff level). The results obtained at these three sites with these controls demonstrated 100% agreement with each other.

Specificity

The following table lists compounds that are detected by the AccuSign™ DOA 4 test. The specificity of the AccuSign™ DOA 4 test was determined by adding various drugs and drug metabolites to drug-negative urine specimens and testing with the AccuSign™ DOA 4 test kit. The results are expressed in terms of the concentration required to produce a positive result. (Table 3.)

Table 3. Specificity

Compound	Concentration (ng/mL)	% Cross-reactivity
THC	Cannabinol	15,000
	11-nor- Δ^1 -THC-9-COOH	30
	11-nor- Δ^1 -THC-9-COOH	50
	Δ^1 -THC	25,000
	Δ^1 -THC	10,000
OPI	Codeine	300
	Glucuronide	300

Hydrocodone	500	60
Hydromorphone	600	50
Levophanol	5,000	6
Meperidine	80,000	0.4
Morphine	300	100
Morphine-3- β -D-glucuronide	500	60
Nalorphine	1,000	30
Naloxane	100,000	0.3
Norcodeine	60,000	0.5
Oxycodone	20,000	1.5
Oxymorphone	60,000	0.5
Procaine HCl	100,000	0.3
Thebaine	5,000	6
COC		
Benzoylegonine	300	100
Cocaine HCl	500	60
Egonine HCl	1,000	30
AMP		
D-Amphetamine	1,000	100
L-Amphetamine	7,000	14
D,L-Amphetamine sulfate	1,000	100
p-OH-Methamphetamine	30,000	3.3
Methylenedioxymphetamine	500	200
Methylenedioxymethamphetamine	10,000	10
β -Phenethylamine	20,000	5
Phentermine	5,000	20
Tryptamine	100,000	1
3-OH-Tyramine	90,000	1.1

The following compounds show no cross-reactivity when tested with AccuSign™ DOA 4 at a concentration of 100 μ g/mL. (Table 4.)

Table 4. Non Cross-Reacting Compounds

Acetaminophen	Dextropropoxyphene	Naproxen
Acetylsalicylate	Diazepam	Norethindrone
Aminopyrine	Diphenylhydantoin	Penicillin
Amitriptyline	Epinephrine	Pentobarbital
Amobarbital	Erythromycin	Phencyclidine
Amoxapine	Estradiol	Phenolbutazone
Ampicillin	Gentisic acid	Phenylpropanol-
Apomorphine	Glutethimide	amine
Ascorbic acid	Guaiacol glycerol	Prednisone
Atropine	ether	Secobarbital
Benzocaine	Hydrochlorothiazide	Tetracycline
Butabarbital	Imipramine	Tetrahydrozoline
Chlordiazepoxide	Lidocaine	Trifluoperazine
Chlorpheniramine	Methadone	Tryptamine
Chlorpromazine	Methaqualone	Zomepirac
Chloroquine	Methyprylon	

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Patent Pending

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AccuSign™ BAR

New One-Step Barbiturates Test

For *In Vitro* Forensic Use Only

Simple One-Step Immunoassay for the Qualitative Detection of Barbiturates in Urine

PBM

Catalog No.	DOA-206	35 Test Kit
	DOA-206-10	10 Test Kit

Intended Use

The AccuSign™ BAR test is a simple, one-step, immuno-chromatographic assay for the rapid, qualitative detection of barbiturates in urine with a cutoff at 300 ng/mL for secobarbital.

The AccuSign™ BAR test provides only a preliminary analytical result. A more specific alternative chemical method must be used in order to obtain a confirmed analytical result. Gas chromatography, mass spectrometry (GC/MS) is the preferred confirmatory method. Other chemical confirmatory methods are available. Clinical consideration and professional judgment should be applied to any drug of abuse test result, particularly when preliminary positive results are used.

Summary and Principle of Procedure

Barbiturates are a group of chemicals derived from barbituric acid. Classified as hypnotics, they depress the central nervous system. Taken orally in pill or tablet form, they are prescribed for many medical conditions, usually for their sedative effect. Abuse of barbiturates can, however, lead not only to impaired motor coordination and mental disorder, but also to respiratory collapse, coma and death. The combination of barbiturates and alcohol is particularly dangerous.

Symptoms of barbiturate abuse include drowsiness, slurred speech and irritability. Acute conditions include respiratory collapse and loss of consciousness. Chronic conditions include addiction, abstinence, seizures, and death. The effects last 3 to 6 hours (10 to 20 hours for phenobarbital). Barbiturates normally remain detectable in urine for 4 to 7 days (up to 30 days for phenobarbital).

Principle

The AccuSign™ BAR test uses solid-phase immunoassay technology for the qualitative detection of secobarbital and barbituric metabolites in human urine. The test is based on the principle of the highly specific immunochemical reactions between antigens and antibodies which are used for the analysis of specific substances in biological fluids. The test relies on the competition for binding to the antibodies between drug conjugates and free which may be present in the urine sample. In the test procedure, a sample of urine is placed in the sample well of the device. It is allowed to migrate upward. If drug is present in the urine sample, it competes with the drug conjugate, which is immobilized on the membrane, for the limited antibodies present in the form of dye-antibody conjugate. When a sufficient amount of drug or drug metabolite above the cutoff level is present, the drug will saturate the antibodies, thus inhibiting the binding of dye-antibody conjugate to the drug conjugate on the membrane. This prevents the formation of a line on the membrane. Therefore, a drug-positive urine sample will not generate a line in the test window, indicating a positive result from positive drug competition, while a negative urine sample will generate a line in the test window, indicating a negative result from an absence of competition with free drug.

In addition to the Test line that may appear in the Test window (T), a Control line is present in the Control window (C) to confirm the viability of the test. This Control line should always be seen if the test is conducted properly. It works as a procedural control, confirming that proper sample volume was used and the reagent system worked. If insufficient sample volume is used, there may not be a Control line indicating the test is invalid.

Materials Provided

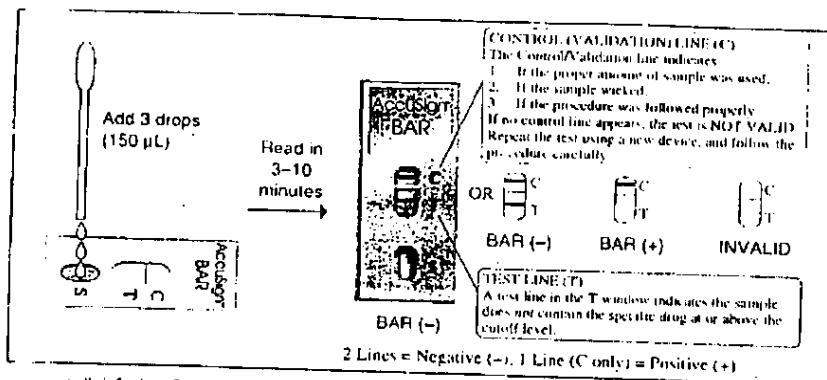
The AccuSign™ BAR test kit contains all the reagents necessary to perform the assay.

- AccuSign™ BAR device. The test device contains a membrane coated with drug conjugate and a pad containing polyclonal anti-barbiturate antibody-alive conjugate in a protein matrix.
- Disposable sample dispenser.
- Instructions for use.

Precautions

- For *in vitro* forensic use only.
- Avoid cross-contamination of urine samples by using a new urine specimen container and dropper for each urine sample.
- This test kit does not contain any HIV or hepatitis infective components. However, urine specimens

EXHIBIT 2



potentially infectious. Proper handling and disposal methods should be followed, according to good laboratory practices.

- The AccuSign™ device should remain in its original sealed pouch until ready for use.
- Do not use the test kit after the expiration date.

Storage and Stability

The AccuSign™ BAR test kit should be stored at 2–30°C (36–86°F) in the original sealed pouch. The expiration date was established under these storage conditions.

Specimen Collection and Preparation

Approximately 150 μL of urine sample is required for each test. Fresh urine specimens do not require any special handling or pretreatment. Specimens should be collected in a clean glass or plastic container. If testing will not be performed immediately, specimens should be refrigerated (2–8°C) or frozen. Specimens should be brought to room temperature before testing.

Specimens containing a large amount of particulate matter may give inconsistent test results. Such specimens should be clarified by centrifuging or allowing to settle before testing.

Test Procedure

The test procedure consists of adding the urine sample to the Sample well of the device and watching for the appearance of colored lines in the result window.

Test Protocol

1. For each test, open one AccuSign™ BAR pouch and label the AccuSign™ device with the patient ID.
2. Holding the dropper vertically, dispense 3 drops (150 μL) of the urine sample into the Sample well (S).
3. Read the result after 3 minutes, but within 10 minutes of sample addition.

Interpretation of Results

Negative: Two Lines. The appearance of two reddish-purple lines—one in the Test window (T) and the other in the Control window (C)—indicates a negative test result (i.e., no barbiturates above the cutoff level have been detected. The color intensity of the Test line may be weaker or stronger than that of the Control line. A negative test result does not necessarily indicate the absence of drug in the sample; it only indicates the sample does not contain drug above the cutoff level in qualitative terms.

Positive: One Line. The appearance of only one reddish-purple line in the Control window (C) and no distinct line in the Test window (T) indicates the test result is positive (i.e., the specimen contains barbiturates at a concentration above the cutoff level). A positive test result does not provide any indication of the level of intoxication or urinary concentration of the drug in the sample; it only indicates the sample contains drug above the cutoff level in qualitative terms.

Invalid: A distinct colored line should always appear in the Control window (C). The test is invalid if no line forms in the Control window (C). Such tests should be repeated with a new AccuSign™ BAR test device.

Note: A very faint line in the Test window (T), visible in 10 minutes, indicates that the amount of barbiturates in the sample is near or below the cutoff level of the test. These urine specimens must be retested, or confirmed with a more specific alternative method such as gas chromatography/mass spectrometry, before positive determinations are made.

Limitations

- The test is designed for use with unadulterated urine only.
- There is a possibility that factors such as technical or procedural errors, as well as other substances in the urine sample which are not listed in Tables 2 or 3 below, may interfere with the test and cause erroneous results.
- Adulterants, such as bleach and/or alum, in urine specimens may produce erroneous results regardless of the method of analysis. If adulteration is suspected, the test should be repeated with a new sample.
- This test detects only the presence of barbiturates and/or their derivatives in urine. A positive test result does not provide any indication of the level of intoxication or urinary concentration.
- The test result read after 10 minutes may not be consistent with the original reading obtained within the 10 minute reading period. The test must be read within 10 minutes of sample application.
- Certain medications containing barbiturates may produce a positive result in any chemical or immunological assay.

User Quality Control

Quality Control: Control standards are not supplied with this kit; however, it is recommended that a control be tested in good laboratory testing practice. NIDA recommends that positive quality control specimens be at or near the cutoff concentration. For information on how to obtain controls, contact PBM's Technical Services. Before using a new kit with patient specimens, positive and negative controls should be tested to confirm the test procedure, and to verify the tests produce the expected Q.C. results. Q.C. specimens should also be run anytime there is any question concerning the validity of results obtained.

Process Control: The Control line can be considered an internal process control. A distinct reddish-purple Control line should always appear if the test procedure is performed properly, an adequate sample volume is used, the sample and reagent are wicking on the membrane, and the test reagents are working. If the Control line does not appear in the control or validation line area, the test is invalid and a new test should be performed. If the problem persists, contact PBM for technical assistance.

Exposure Values

AccuSign™ BAR is a qualitative assay. The amount of secobarbital or barbiturate metabolites present in the urine cannot be estimated by the assay. The assay results distinguish positive from negative samples. Positive results indicate the samples contain barbiturates above the cutoff concentration.

Performance Characteristics

The AccuSign™ BAR test has been shown to detect an average of 300 ng/ml of secobarbital in urine. The test also detects other barbiturates listed below at the minimum concentrations indicated (Table 2).

The accuracy of AccuSign™ BAR was evaluated in comparison to a commercially available immunoassay (Syva®EMIT® II). A total of 302 samples was tested by both procedures. The overall accuracy of the test was 98.7%, as shown below (Table 1.)

Table 1. Accuracy: Comparison of AccuSign™ BAR with Syva®EMIT® II

Syva®EMIT® II (BAR)			
	Positive	Negative	TOTAL
AccuSign™	105	0	105
BAR	4	193	197
TOTAL	109	193	302

Relative Sensitivity: AccuSign™ BAR = 96.3% (105/109) > 99.9% (193/193)

Precision and Accuracy

The precision of AccuSign™ BAR was determined by carrying out the test with serially diluted standard drug solutions. About 98% of the samples containing drug levels 25% over the cutoff level consistently showed positive results.

The study also included over 40 samples \pm 25% cutoff level as a challenge of cutoff precision. These results were found to be consistently in agreement with expected test results.

Distribution of Random Error:

Twenty (20) blind samples prepared by spiking various concentrations of drug were separately tested by two operators. The test results from the two operators showed complete agreement.

Reproducibility

The reproducibility of the test results of AccuSign™ BAR was examined at three different sites using a total of 15 blind controls, consisting of 5 negative samples, 5 moderately positive samples, and 5 strongly positive samples (i.e., a concentration 3 times the cutoff level). The results obtained at these three sites with these controls demonstrated 100% agreement with each other.

Specificity

Compounds that are detected by the AccuSign™ BAR test are listed below (Table 2). The specificity of AccuSign™ BAR was determined by adding various drugs and drug metabolites to drug-negative urine specimens and testing with the AccuSign™ BAR test kit. The results are expressed in terms of the concentration required to produce a positive result.

Table 2. Specificity

Compound	Concentration (ng/mL)	% Cross-reactivity
Allobarbital	200	150
Alphenal	1,000	30
Aminobarbital	2,000	15
Apdrobarital	200	150
Barbital	2,000	15
Butabarbital	500	60
Butalbital	200	150
Cyclopentobarbital	500	60
Pentobarbital	1,000	30
Phenobarbital	5,000	0
Secobarbital	300	100

The following compounds show no cross-reactivity when tested with AccuSign™ BAR at a concentration of 100 µg/mL. (Table 3)

Table 3. Non Cross-Reacting Compounds

4-Acetamidophenoxy	Chlorpromazine	(+)-Epinephrine
Acetophenetidin (Phenacetin)	Chlorquine	(-)-Epinephrine
N-Acetylprocainamide	Cholesteryl	(-)- ¹ H-Epinephrine
Acetyl Salicylic acid	Chlorpromazine	Brylaurysyl
Amphetamine	Chlorzine	B-Galactosidase
Amphetamine	Clomipramine	Estone-19-carboxylic
Amphetamine	Codeine	Ethyldiisopropyl-
Amphetamine	Coramine	butyrate
Amphetamine	Cortisone	Fenoprofen
D,L-Amphetamine	Creatinine	Flavonoids
L-Amphetamine	Desmethylcorticososterone	Gastric acid
Amorphophane	Desmethylmorphinan	Glucuronic acid
Aspartame	Desoxepin	Glutathione
Atropine	Desoxynorepinephrine	Guaiacol
Benzoic Acid	Desoxytubocurarine	Hippurate
Benzene and	Desmethylatropine	Hydralazine
Benzylcyclohexene	Desmethylcortisol	Hydrochlorothiazide
Benzylcyclohexene	Desmethylcortisone	Hydrocodone
Cannabidiol	Desmethylcortisone	Hydrocortisone
Clibutidine	Desmethylcortisol	Hydroxyzine
Chlordiazepoxide	Desoxynorepinephrine	Hydromorphone
Chlordiazepoxide	Desoxynorepinephrine	O-Hydroxyhippuric
Chlorpromazine	Desoxynorepinephrine	acid
Chlorpromazine	Desoxynorepinephrine	β-Hydroxytyramine
Chlorpromazine	Desoxynorepinephrine	

Ibuprofen	Naproxen	Primidone
Isoproterenol	Nifedipine	Propiophenone
(+)-Isoproterenol	Nicodine	D,L-Propiopanol
Isosorbide	Nimodipine	Propiobutanone
Ketamine	Nitrosoxymephedrone	D-Pipecolophenone
Ketoprofen	D-Norpseudoephedrine	D-Pipecolylethylene
Labetalol	(-)-Norpseudo-	Quindine
Lorazepam	eophedrine	Quinine
Lubagaine	Nisoldipine	Ranitidine
Loperamide	Oxazine	Salicylic acid
Loratadine	O-D,L-Oxipipame	Saxenda
Luzopine succinate	Oxalic acid	Serotonin
Meprobamate	Oxazepam	Sulfamethazine
Meprobamate	Oxazoline Acid	Sulindac
Methadone	Oxazolidine	Tetrahydrocannabinol
p-Hydroxymethyl-	Oxymecatetra-	Tetrahydrocannabinol
-phenacetin	Phenacetin	Theanine
Methaqualone	Papaverine	Thiamine
Methoxyphenamine	Paracetamol	Thiobutazine
(D)-2-Methylc-	Paracetamol	D,L-Thymine
-dihydromamphe-	Paracetamol	Tolbutamide
-amine	Paracetamol	Tramadol
(L)-2-Methylc-	Paracetamol	Trifluoperazine
-dihydromamphe-	Paracetamolazine	Trimeptopan
-amine	Paracetamol	Trimipramine
Methylphenidate	Paracetamol	Tryptamine
Methyprylon	Phenacetin	D,L-Tryptophan
Morphine-3-O-D-	Phenacetin	Tryptamine
-glucuronide	Phenylethylamine	D,L-Tryptophan
Naloxone and	Phenylpropylamine	Tryptamine
Melatonin	Phenylpropylamine	D,L-Tryptophan
Naloxone	Phenylpropanolamine	Uric acid
Naloxone	Phenyltoluenesulfonate	Vigaparin
Naloxone	Phenyltoluenesulfonate	Zolmetopan

References

- Hawks RL, Chang CN, eds. *Urine Testing for Drugs of Abuse*. Rockville, MD: National Institute on Drug Abuse (NIDA); Research Monograph 73, 1996.
- Baech RC. *Disposition of Oral Drugs and Chemicals in Man*. 2nd Ed. Davis, CA: Brodman; 1982, p 468.

PBM

Accusign™ is a Trademark of Princeton BioMedTech Corporation.
Patent Pending

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96A 19 02/2002

Princeton BioMedTech Corporation
P.O. Box 7139, Princeton, New Jersey 08543-7139 USA
1212 U.S. Route 1, Monroe, New Jersey 08831 USA
Tel: (609) 274-1000
Fax: (609) 274-1010

4



Presents the AccuSign™ DOA Series
**One-Step Drug Test with
Results in Only 2-5 Minutes**



- Easy to Read Color
- Highly Sensitive
- Built-In Test Control
- No Refrigeration
- Eliminates Timing
- Simultaneous Testing Capability of Multi-Drugs

Tests are available in single or multiple test panels for the following drugs:

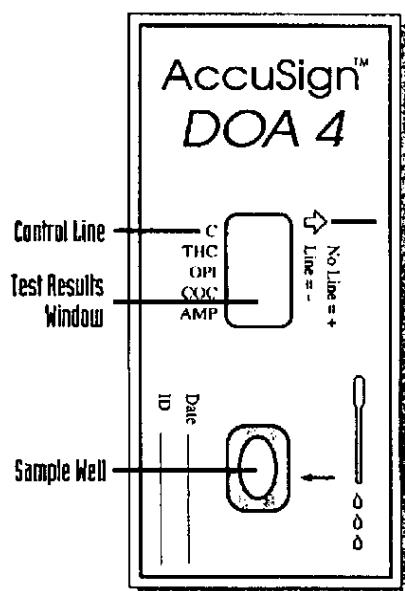
Amphetamines	Cocaine	PCP
Barbiturates	Methamphetamine	THC [Marijuana]
Benzodiazepines	Morphine/Opiates	

Watch us for More Tests & New Technology

EXHIBIT 3

AccuSign™ DOA Series

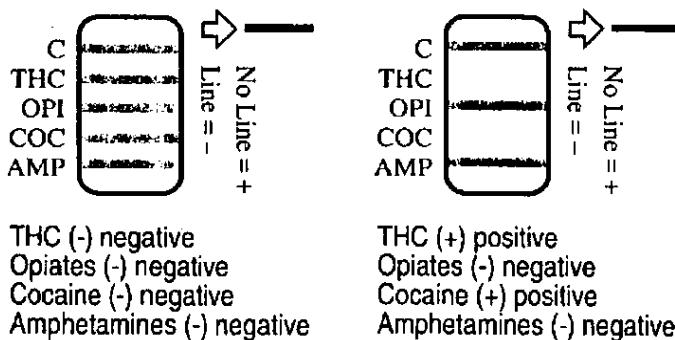
Test Procedure



1. Using the plastic pipette, add 3 drops of urine sample to the Sample Well.
2. Read results in 2 to 5 minutes [within 10 minutes].
3. Interpret Test Results Window:
 - **CONTROL LINE** - A colored line indicates the test is complete and the system has worked properly.
 - **NEGATIVE** - A colored line for the specific drug indicates the test is negative and the drug was **NOT DETECTED**.
 - **POSITIVE** - No colored line for the specific drug indicates the test is positive and the drug was **DETECTED**.

Tests Manufactured by Princeton BioMeditech Corporation

Samples



For information or to place an order call:

UNIVERSAL DRUG TESTING
478 ROUTE 61
LARGE, PA 18025

Visualine™ II

One Step Drug Screening Test

AVITAR TECHNOLOGIES, INC.
produces, markets and distributes medical devices for
the health care industry.

Avitar now distributes rapid Tests for Drug Abuse
that are:

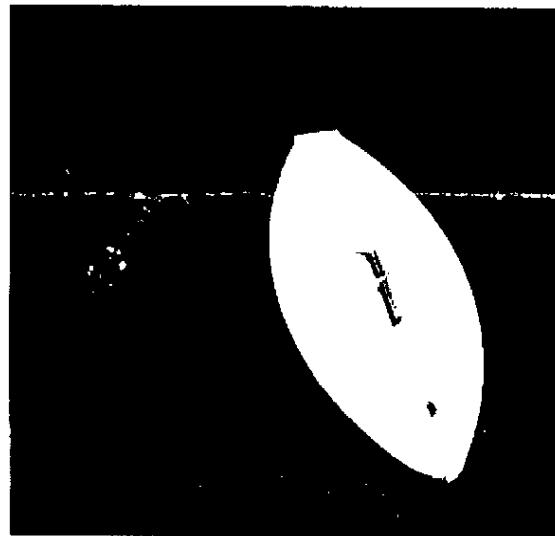
- accurate
- sensitive
- simple to use

The Visualine™ II tests¹ which have FDA approval to
market are:

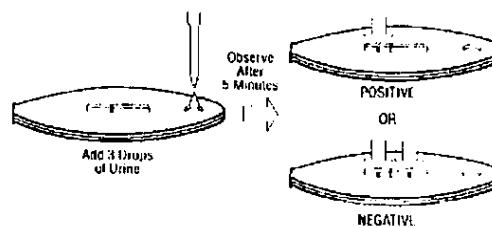
- Cocaine
- Morphine
- Cannabinoids (THC)
- Benzodiazepines

The tests are composed of preformulated dry reagents
arranged on a porous membrane support in a convenient
cassette. To perform the tests, just add a few drops of urine to
the sample well of the device and wait five minutes. Results
are then easily read in the results window as the presence or
absence of a red line. A built in reference control ensures that
the sample has been added and that the test is effective. The
tests are based on the newest lateral flow micro-particle
immunoassay technology and have been evaluated in clinical
trials at a major university. The Visualine™ II tests are
designed to meet NIDA proposed cutoff levels.

¹ A test for Methamphetamine is available for uses that do not require FDA
approval to market.



Visualine™ II test device and pipette.



Simple to use and simple to read.

For information contact customer service at
1.800.255.0511

AVITAR TECHNOLOGIES, INC.
65 Dan Road, Canton, Massachusetts 02021

EXHIBIT 4

HOME DRUG TEST

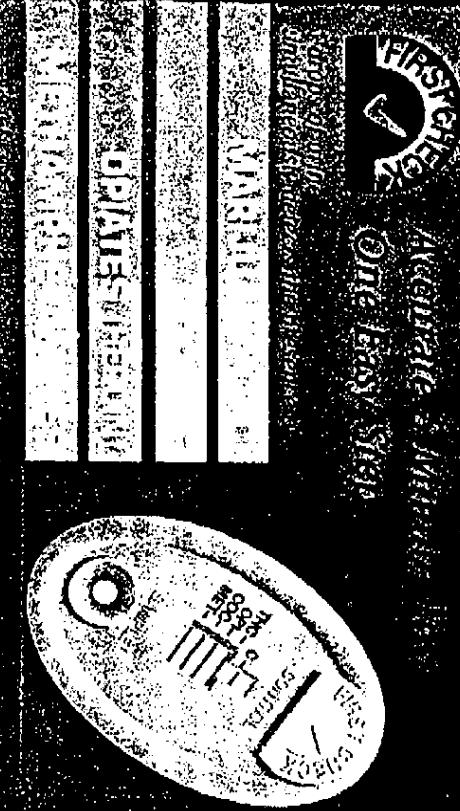
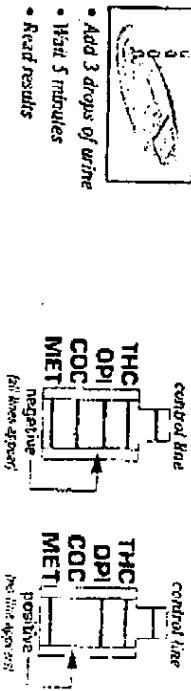


EXHIBIT 5

For information about other home drug tests call 888-788-5716.
Worldwide Medical Corporation, Irvine, CA 92618 • <http://www.wmmed.com>

EASY TO TEST**EASY TO READ RESULTS**

First Check² provides immediate information about the use of marijuana, cocaine, opiates and methamphetamine. It is not for legal, law enforcement, or medical purposes. For diagnosis and treatment, consult with health care or substance abuse professional.

This box contains:

Material required but not provided:

- I test well
- I sample digester
- I collection of sample.

Instructions

For external use only. Store at 40-65°F [4-30°C]. Read enclosed directions completely before use.

Current Drug Usage Trends:

Marijuana use increased 43% in junior high (grades 6, 8) and 28% in high school students (grades 9-12).

National Survey's "Resource Institute for Drug Education, Inc. (F.R.I.D.E.) 9th annual students' survey (September 25, 1995)

In 1995, cocaine related episodes comprised 27% of all emergency department drug related episodes.

1995 Preliminary Estimates of Drug Related Emergency Department Episodes, SAMHSA/CMS (August, 1996)

The potency of marijuana has doubled since the 1970s.
1995 Preliminary Estimates of Drug Related Emergency Department Episodes, SAMHSA/CMS (August, 1996)
Marijuana Research Project Quarterly Report, 1996, Mt. Sinai: University of Mississippi, Research Institute of Pharmaceutical Sciences

Methamphetamine-related emergency department episodes rose 25% between 1991 and 1994.

1995 Preliminary Estimates of Drug Related Emergency Department Episodes, SAMHSA/CMS (August, 1996)

Among emergency room cocaine-related episodes, "dependence" was the most commonly reported motive for drug use in 1995.

1995 Preliminary Estimates of Drug Related Emergency Department Episodes, SAMHSA/CMS (August, 1996)

In 1994, which manner of drug abuse death was accidental, cocaine was mentioned in 55% and heroin/morphine in 51% of medical examiner cases.

Statistical Series I, Number 14-B, Drug Abuse Warning Network (DAWN), U.S. Dept. of Health and Human Services

A higher prevalence of depression, motivational problems, and interpersonal problems are associated with marijuana use.

Corus Foundation, Marijuana and Today's Youth (1997)

Date: 5/26/04 Time: 2:45:00 PM

From: Sullivan To: Extr. Lyle Alexander

WORLDWIDE MEDICAL

CO. P. O. R. A. T. O. N.

harm, U.S.A. 92616 U.S.A.

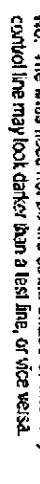
Call: 800-778-5216

Visit Worldwide Medical Corporation at <http://www.wm.com>

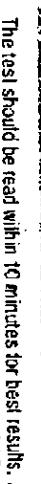
Commonly Asked Questions:

G. Test Results

A: No. The lines need not be the same shade or intensity. The control line may look darker than a test line, or vice versa.

C: 

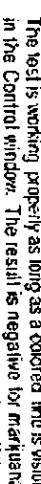
A: The test should be read within 10 minutes for best results. A negative result (5 lines, 1 line in Control window and 4 lines in Test window) will never disappear.

C: 

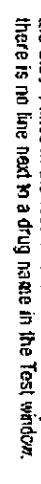
A: The test is working properly as long as a colored line is visible in the Control window. The result is negative for marijuana, morphine/opiates, cocaine and methamphetamine if there are also 4 lines in the Test window. The result is positive if there is no line next to a drug name in the Test window.

C: 

A: Yes, we recommend that you wait the full 5 minutes before reading the result. However, if your test clearly shows negative (5 lines, a line in the Control window and 4 lines in the Test window), you do not need to wait. If a line in the Test window is not clear, test again. Your result will be more reliable after 5 minutes. Most negative test results will be more reliable in 1 minute. To be sure of a positive (1 line in Control window and no line next to a drug name in the Test window), wait 5 minutes but no longer than 10 minutes.

O: 

A: Parentally gain your child's cooperation with the understanding that it is their welfare that you are protecting. Should your child question the importance of testing him or her for drugs or the seriousness of occasional drug usage, page 4 of this pamphlet supplies current data on drug usage among our youth and why your child cannot ignore this possibility. Be persistent without accusations, threats, or anger. If your child continues to refuse, seek professional help. Check with your physician, your phone directory also provides sources of assistance under "Alcohol and Drug Abuse" in the Government and Community Services Listings, or page 4 of this pamphlet for information by Services Listings, or page 4 of this pamphlet for more information.

O: 

A: Even though the result is negative, I still feel that my child may be using drugs. What can I do?

O: Check with your physician, your phone directory also provides sources of assistance under "Alcohol and Drug Abuse" in the Government and Community Services Listings, or page 4 of this pamphlet for Support Group information.

HOME DRUG TEST
Instructions for Use



First Check[®]
Marijuana (THC),
Morphine/Opiates,
Cocaine, &
Methamphetamine

When the need to know is...now

- Simple - one step

• Easy-to-read

• Confidential

• Result in 5 minutes

First Check[®] Marijuana, Morphine/Opiates, Cocaine, & Methamphetamine provides immediate information about the use of marijuana, morphine/opiates, cocaine, and methamphetamine. It is useful for legal, law enforcement, or medical purposes. For diagnosis and treatment, consult with a healthcare or substance abuse professional.

Read the following directions completely before use.

FOR EDUCATIONAL USE ONLY
Store at 40-86°F (4-30°C).
For external use only.

Store at 40-86°F (4-30°C).

First Check® Marijuana (THC), Morphine/Opiates, Cocaine, & Methamphetamine

Not to be taken internally.

Marijuana
THC is the primary active ingredient in marijuana (cannabinoids). When ingested or smoked, it produces euphoric effects. Users have impairment of short term memory and marijuana use slows learning. Also, it may cause transient episodes of confusion, anxiety, or even frank toxic delirium. Long-term, relatively heavy use may be associated with behavioral disorders. The peak effect of smoking marijuana occurs in 20-30 minutes and the duration is 90-120 minutes after one cigarette. Elevated levels of urinary metabolites are found within hours of exposure and remain detectable for 3-10 days after smoking.

Morphine/Opiates

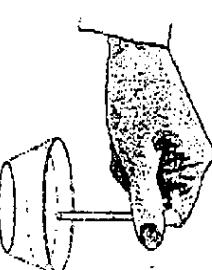
Opioid analgesics comprise a large group of substances which control pain by depressing the central nervous system. Morphine is the prototypical compound of this group. Up to 75% of the morphine dose is eliminated in the urine as glucuronide. Free morphine in the urine accounts for about 11% of the dose, while very small amounts of morphine-6-glucuronide and morphine-3-sulfate sulfate are also present. Approximately 5% of a dose of morphine is N-deacetylated to normorphine, which is found as a urinary metabolite in both free (15%) and conjugated (45%) forms. Cocaine is excreted in the urine primarily as amphetamine and oxidized and deaminated derivatives, cocaine, and methamphetamine. However, 10-20% of methamphetamine is excreted unchanged. Thus the presence of the parent compound in the urine indicates methamphetamine use. Methamphetamine is generally detectable in the urine for 3-5 days, depending on urine pH level.

Before you begin

Read all the information in this pamphlet before performing the test. First, make sure you are familiar with the test kit contents shown below. Store at 16-36°C (2-20°C) in the sealed pouch, away from direct sunlight. Do not use after the expiration date stamped on the package.

Instructions

1. Open the sealed pouch, remove the First Check® card, and set the card on a flat surface with Test and Control windows facing up.
2. Collect urine sample in a clean plastic or glass container.



Limitations

The First Check® One-Step Home Drug Test is not reusable. The test instructions must be followed precisely.

The test detects only the presence of marijuana (THC), morphine, cocaine, and methamphetamine or their metabolites in urine. A positive test does not provide any information about the amount or level of intoxication.

The test is designed for use with undiluted urine only. Adults, certain salts, such as bleach and/or alum, in a urine sample may produce an erroneous result. If adulteration is suspected, the test should be repeated with a new urine sample.

The result must be read 5-10 minutes after sample application. A result read after 10 minutes may not be accurate.

Urine sample should be at room temperature. If sample has been refrigerated, allow sample to reach room temperature before testing.

Certain medications containing opiates or methamphetamine may produce a positive result in any chemical and immunological assay. Autoradiographs and tea containing poppy products and/or tea leaves may produce a positive result. Prolonged exposure to second-hand marijuana smoke may produce a positive result.

Methamphetamine

Methamphetamine is a potent sympathomimetic agent with therapeutic applications. The drug can be taken orally, injected, or inhaled. Acute higher doses lead to enhanced stimulation of the central nervous system and induce euphoria. Cardiovascular responses to methamphetamine include increased blood pressure and cardiac arrhythmia. More acute responses include anxiety, psychosis, hallucinations, psychiatric behavior, and euphoria, depression, and exhaustion. The effects of methamphetamine generally last 2-4 hours, and the drug has a half-life of 9-14 hours in the body. Methamphetamine is excreted in the urine primarily as amphetamine and oxidized and deaminated derivatives. However, 10-20% of methamphetamine is excreted unchanged. Thus the presence of the parent compound in the urine indicates methamphetamine use. Methamphetamine is generally detectable in the urine for 3-5 days, depending on urine pH level.

Results

Wait at least 5 minutes but not more than 10 minutes before reading result.

Negative

Five horizontal lines, one line in the Control (upper) window and four lines in the Test (lower) window, mean there is no marijuana, morphine, cocaine, nor methamphetamine present in the urine sample. The fine line in the Test window may be lighter or darker than the line in the Control window.

No drug taken

Positive
(one line in the Control window and no line next to a drug name in the Test window means the sample contains that drug)
Drug Taken (Cocaine)

A distinct colored line should always appear in the Control (upper) window. If no line appears in the Control window, do not interpret result.

Invalid Test

The test is designed for use with undiluted urine only. Adults, certain salts, such as bleach and/or alum, in a urine sample may produce an erroneous result. If adulteration is suspected, the test should be repeated with a new urine sample.

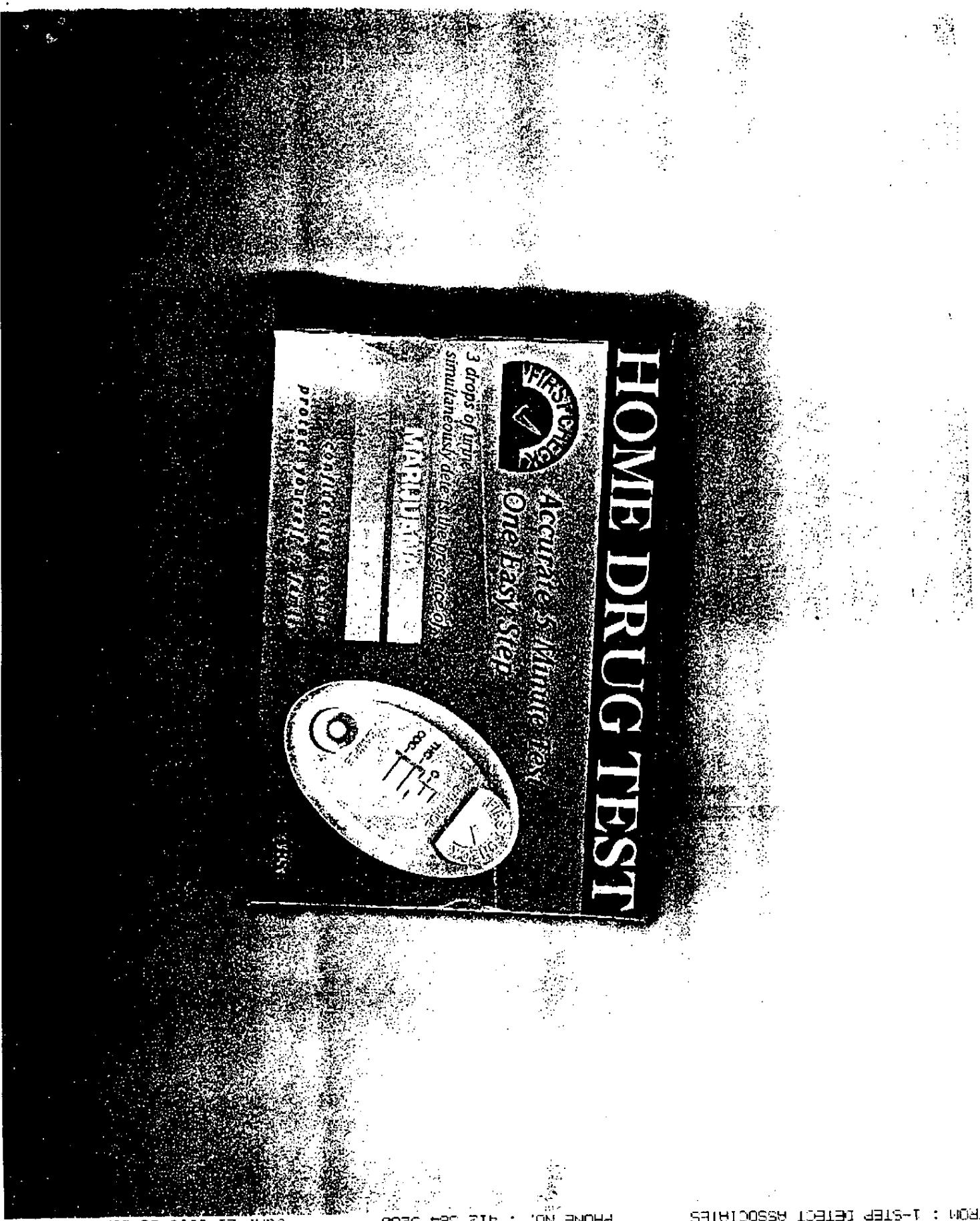
The result must be read 5-10 minutes after sample application. A result read after 10 minutes may not be accurate.

Urine sample should be at room temperature. If sample has been refrigerated, allow sample to reach room temperature before testing.

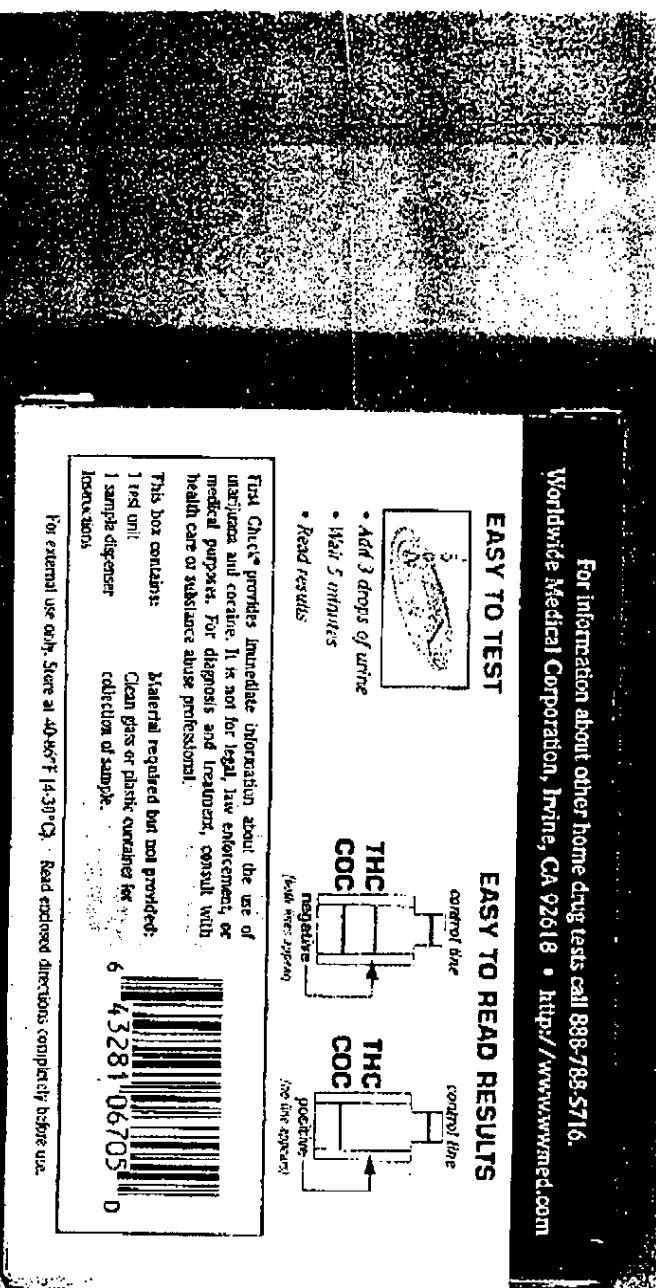
Certain medications containing opiates or methamphetamine may produce a positive result in any chemical and immunological assay.

Autoradiographs and tea containing poppy products and/or tea leaves may produce a positive result. Prolonged exposure to second-hand marijuana smoke may produce a positive result.

Control	Glass plastic or glass container
Water	Water for sample collection
Test	Test Window
Wash	not provided
Sample	Sample Dispenser
First Check®	



FROM : 1-STEP DETECT ASSOCIATES PHONE NO. : 412 384 5260 Jun. 21 1999 08:35AM PS



From: Sullivan To: Exr. Lyle Alexander

Marijuana use increased 43% in junior high (grades 6-8) and 28% in high school students (grades 9-12).
 National Parent Resource Institute for Drug Information, Inc. [2001]
1995 annual student survey (September 21, 1996)
 Epilepsy, Substance Abuse (August, 1996)
 The potency of marijuana has doubled since the 1970s.
Marijuana Potency Monitoring Project: Quarterly Report, 1996, Mississippi University of Mississippi, Research Institute of Pharmacology and Sciences

Heavy marijuana use is associated with reduced ability to sustain attention, a decreased capacity to shift attention, reduced learning and decreased mental flexibility.

Pope H. and Henklein, Test 11, Journal of the American Medical Association, 273, 523-527 (1995).

Among emergency room cocaine-related episodes, "dependence" was the most commonly reported motive for drug use in 1995.
1995 Preliminary Estimates of Drug-Related Emergency Department Episodes, SAMHSA/CDAS (August, 1996).

Educational Materials

In 1994, when number of drug abuse death was accidental, same was mentioned in 55% of medical examiner cases.
Statistical Series, Series I, Number 14-2, Drug Abuse Warning Network (DAWN), U.S. Dept. of Health and Human Services

A higher prevalence of depression, motivational problems and interpersonal problems are associated with marijuana use.
Caren Foundation, Marijuana and Today's Youth (1997).

Date: 5/26/04 Time: 2:45:00 PM

Commonly Asked Questions:

HOME DRUG TEST

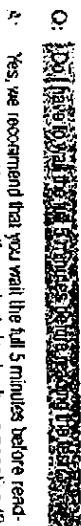
Instructions for Use



First Check® Marijuana & Cocaine

When the need to know is...now

- Simple - one step



C:

D:

O:

A:

The test is working properly as long as a colored line is visible in the Control window. The result is negative for marijuana and cocaine. If there are also 2 lines in the Test window, the result is positive if there is no line next to a drug name in the Test window.

- Result in 5 minutes

First Check® Marijuana & Cocaine provides immediate information about the use of marijuana and cocaine. It is not for legal, law enforcement, or medical purposes. For diagnosis and treatment, consult with a health care substance abuse professional.

• Easy-to-read

• Confidential

Read the following directions completely before use.

- Save the unused portion of the urine sample. Ask your family physician for a recommended laboratory to retest the same sample.

For external use only.

Store at 36-86°F (2-30°C).



Visit Worldwide Medical Corporation at <http://www.worldmed.com>



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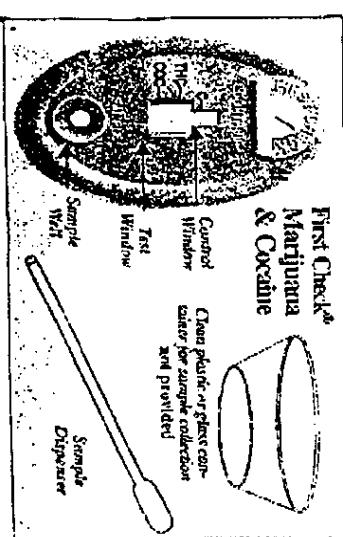
MM:

NN:

First Check®

Marijuana & Cocaine

Not to be taken internally.



Date: 5/26/04 Time: 2:45:00 PM

Marijuana & Cocaine

THC is the primary active ingredient in marijuana (cannabinoids). When ingested or smoked, it produces euphoric effects. Users have impairment of short term memory and marijuana use slows learning. Also, it may cause transient episodes of confusion, anxiety, or even frank toxic delirium. Long term, relatively heavy use may be associated with behavioral disorders. The peak effect of smoking marijuana occurs in 20-30 minutes and the duration is 90-120 minutes after one cigarette. Elevated levels of urinary metabolites are found within hours of exposure and remain detectable for 3-10 days after smoking.

Cocaine derived from the leaves of coca plant is a potent central nervous system (CNS) stimulant and a local anesthetic. Cocaine induces euphoria, confidence and a sense of increased energy in the user; these psychological effects are accompanied by increased heart rate, dilation of the pupils, fever, tremors and sweating. Cocaine is used by smoking, intravenous, intranasal or oral administration and excreted in the urine primarily as benzoylecgonine in a short time. Benzoylecgonine

has a longer biological half-life (3-6 hours) than cocaine (0.5-1.5 hours) and can generally be detected for 24-60 hours after cocaine use or exposure.

Before you begin

Read all the information in this pamphlet before performing the test. First, make sure you are familiar with the test kit contents shown, below. Store at 36-86° F (2-30° C) in the sealed pouch, away from direct sunlight. Do not use after the expiration date stamped on the package.

Instructions

1. Open the sealed pouch, remove the First Check® card, and set the card on a flat surface with Test and Control windows facing up.

2. Collect urine sample in a clean plastic or glass container.

3. With sample dispenser over sample, press bulb between thumb and index finger; insert dispenser opening into sample and release pressure on bulb. Sample will fill half of dispenser tube.

- Do not discard the unused urine sample after the test has been completed and the result is known.*
4. With sample dispenser in vertical position over Sample well of test card, gently squeeze dispenser bulb. To allow 3 fully-formed drops of urine, one at a time, to fall into Sample well.

Results
A line at least 5 minutes but not more than 10 minutes after reading result.

Negative

Three horizontal lines, one line in the Control (upper) window and two lines in the Test (lower) window, means there is no marijuana and no cocaine present in the urine sample. *The lines in the Test window may be lighter or darker than the line in the Control window.*

No drug taken

Two horizontal lines, one line in the Control (upper) window and one line in the Test (lower) window, means there is no marijuana and no cocaine present in the urine sample. *The lines in the Test window may be lighter or darker than the line in the Control window.*

Positive

One line in the Control window and no line next to a drug name in the Test window means the sample contains that drug.

Drug taken (cocaine)

A distinct colored line should always appear in the Control (upper) window. If no line appears in the Control window, do not interpret result.

Limitations

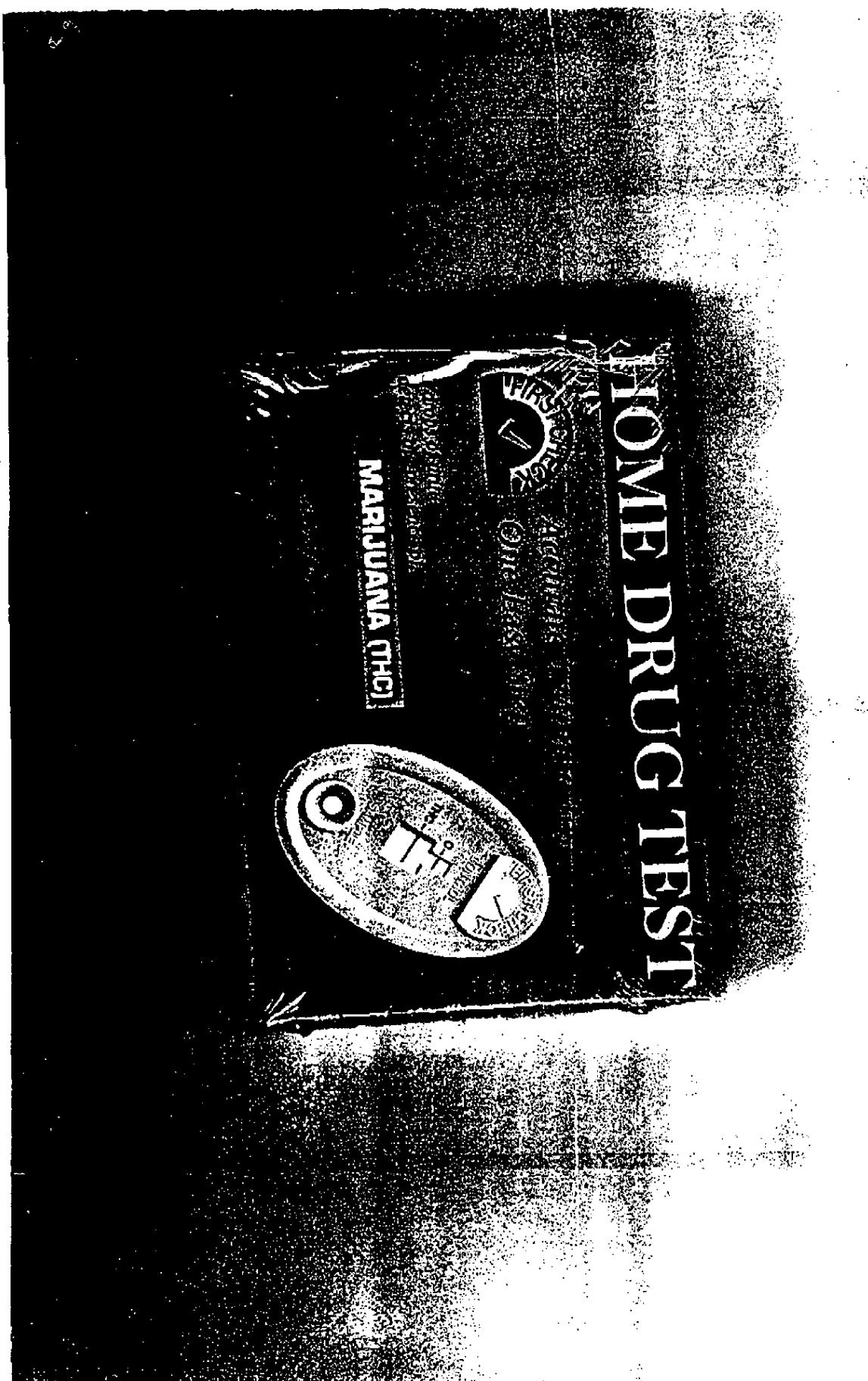
The First Check® One-Step Home Drug Test is *not* reusable. The test instructions must be followed precisely.

The test detects only the presence of marijuana (THC) and cocaine or their metabolites in urine. A positive test does not provide any information about the amount or level of intoxication.

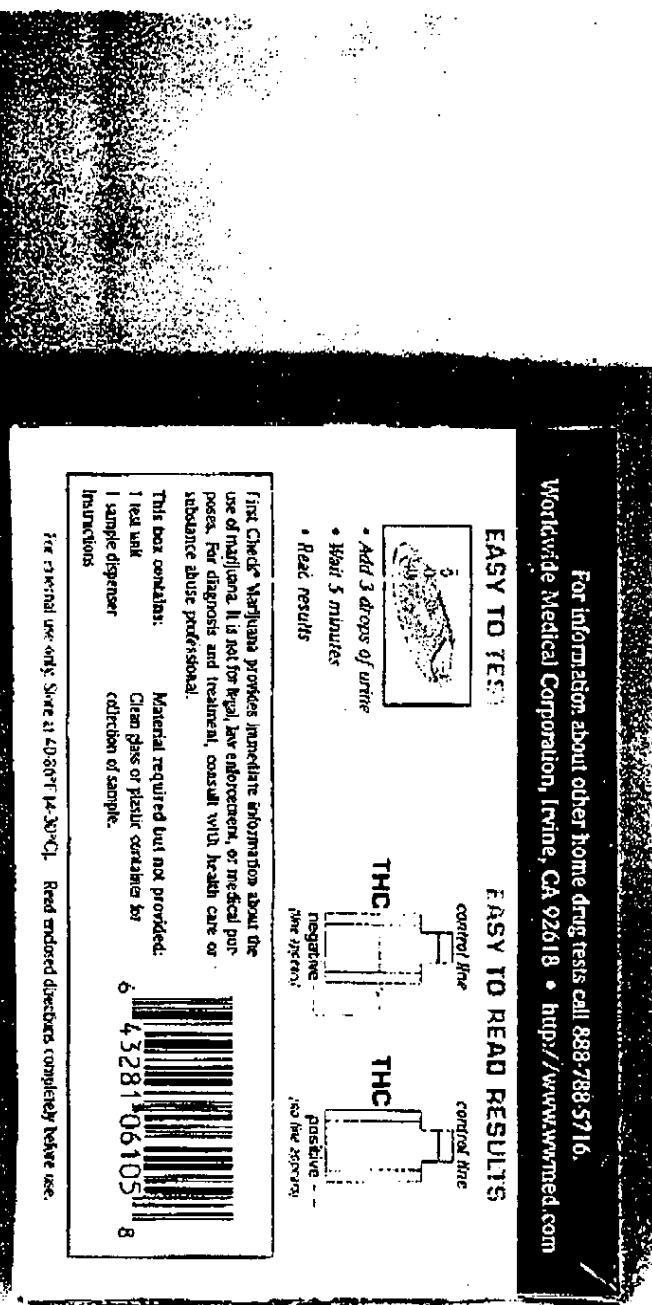
The test is designed for use with undiluted urine, only. Adulterants, such as bleach and/or alum, in a urine sample may produce an erroneous result. If adulteration is suspected, the test should be repeated with a new urine sample. The result must be read 5-10 minutes after sample application. A result read after 10 minutes may not be accurate.

Urine sample should be at room temperature. If sample has been refrigerated, allow sample to come to room temperature before testing.

Prolonged exposure to secondhand marijuana smoke may produce a positive result.



FROM : 1-STEP DETECT ASSOCIATES PHONE NO. : 412 324 5660 Jun. 21 1999 08:42AM P11



First Check® Marijuana provides immediate information about the use of marijuana. It is not for legal law enforcement, or medical purposes. For diagnosis and treatment, consult with health care or substance abuse professional.

This test contains:

- 1 tea bag
- 1 sample dispenser
- Instructions

Material required but not provided:
Clean glass or plastic containers for collection of sample.

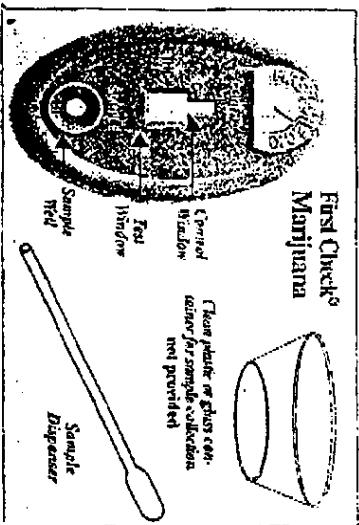
For external use only. Store at 40-80°F (4-30°C). Read enclosed directions completely before use.

Date: 5/26/04 Time: 2:45:00 PM

First Check®[®]

Marijuana (THC)

Not to be taken internally.



Before you begin

Read all the information in this pamphlet before performing the test. First, make sure you are familiar with the test kit contents shown, below. Store at 36-86°F (2-30°C) in the sealed pouch, away from direct sunlight. Do not use after the expiration date stamped on the pack.

Simple & Accurate
Just one step
No need to wait
Read results in 3 minutes
Accurate: Our 99% accurate laboratory studies show...

Marijuana (THC)

THC is the primary active ingredient in marijuana (cannabinoids). When ingested or smoked, it produces euphoric effects. Users have impairment of short term memory and marijuana use slows learning. Also, it may cause transient episodes of confusion, anxiety, or even frank toxic delirium. Long term, relatively heavy use may be associated with behavioral disorders. The peak effect of smoking marijuana occurs in 20-30 minutes and the duration is 90-120 minutes after one cigarette. Elevated levels of urinary metabolites are found within hours of exposure and remain detectable for 3-10 days after smoking.

Instructions

1. Open the sealed pouch, remove the First Check® card, and set the card on a flat surface with Test card, and Control windows facing up.
2. Collect urine sample in a clean plastic or glass container.



3. With sample dispenser over sample, press bulb between thumb and index finger. Insert dispenser opening into sample and release pressure on bulb. You should see sample fill half of dispenser tube.



(Do not discard the urined until after the test has been completed and the results are read.)

4. With sample dispenser in vertical position over Sample well of test card, gently squeeze dispenser bulb to allow 3 fully-formed drops of urine, one at a time, to free-fall into Sample well.



5. Allow the test card to remain undisturbed until result is read. Read the result after 3 minutes but within 10 minutes.

Results
Wait at least 3 minutes but no more than 10 minutes before reading result.

2 lines - negative

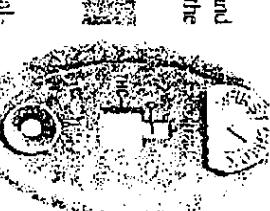


1 line - positive

One line in the Control window and no line in the Test window means the sample contains marijuana.

drug taken

Invalid Test
A distinct colored line should always appear in the Control (upper) window. If no line appears in the Control window, do not interpret result.



Limitations

The First Check® One-Step Home Drug Test is not reusable. The test instructions must be followed precisely.

The test detects only the presence of marijuana (THC) or its metabolites in urine. A positive test (no line in Test window) does not provide any information about the amount or level of intoxication.

The test is designed for use with unadulterated urine, only. Adulterants, such as bleach and/or alum, in a urine sample may produce an erroneous result. If adulteration is suspected, the test should be repeated with a new urine sample.

The result must be read 3-10 minutes after sample application. A result read after 10 minutes may not be accurate.

Urine sample should be at room temperature. If sample has been refrigerated, allow sample to come to room temperature before testing. Prolonged exposure to secondhand marijuana smoke may produce a positive result.